



## United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	NO. CONFIRMATION NO.	
10/814,244	04/01/2004	Steven J. Soldin	31603-2049 6946		
33721 TODAYS LLD	7590 01/02/2008		EXAMINER		
TORYS LLP 79 WELLING	TON ST. WEST	SODERQUIST, ARLEN			
SUITE 3000	NI M5V 1NO	ART UNIT	PAPER NUMBER		
TORONTO, O CANADA	IN IVISK TINZ		1797		
			MAIL DATE	DELIVERY MODE	
			01/02/2008	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

		Applicatio	Application No. Applicant(s)						
		10/814,244	4	SOLDIN, STEVEN J.					
	Office Action Summary	Examiner		Art Unit					
		Arlen Sode	rquist	1797					
	The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply								
WHIC - Exter after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR FOR HEVER IS LONGER, FROM THE MAILIN nsions of time may be available under the provisions of 37 C SIX (6) MONTHS from the mailing date of this communicating operiod for reply is specified above, the maximum statutory or to reply within the set or extended period for reply will, by reply received by the Office later than three months after the ed patent term adjustment. See 37 CFR 1.704(b).	NG DATE OF TH! CFR 1.136(a). In no ever ion. period will apply and will r statute, cause the appli	S COMMUNICATIOnt, however, may a reply be to expire SIX (6) MONTHS from cation to become ABANDON	N. imely filed  n the mailing date of this communication (35 U.S.C. § 133).					
Status									
1)	Responsive to communication(s) filed on				•				
′—	•	 This action is no	n-final.						
/ <del>-</del>	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is								
,—	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.								
Dispositi	ion of Claims								
4)🖂	Claim(s) 1-32 is/are pending in the applic	cation.							
	4a) Of the above claim(s) is/are withdrawn from consideration.								
5)	5) Claim(s) is/are allowed.								
6)⊠	6)⊠ Claim(s) <u>1-32</u> is/are rejected.								
7)	Claim(s) is/are objected to.								
8)□	Claim(s) are subject to restriction	and/or election re	quirement.						
Applicati	ion Papers								
9)[	The specification is objected to by the Exa	aminer.							
10)	The drawing(s) filed on is/are: a)	accepted or b)	$\square$ objected to by the	Examiner.					
	Applicant may not request that any objection	to the drawing(s) be	e held in abeyance. Se	e 37 CFR 1.85(a).					
	Replacement drawing sheet(s) including the o								
11)	The oath or declaration is objected to by t	the Examiner. Not	te the attached Office	e Action or form PTO-152	2.				
Priority (	under 35 U.S.C. § 119								
-	Acknowledgment is made of a claim for fo ☐ All b) ☐ Some * c) ☐ None of:	oreign priority und	er 35 U.S.C. § 119(a	a)-(d) or (f).					
	1. Certified copies of the priority documents have been received.								
	2. Certified copies of the priority documents have been received in Application No								
	3. Copies of the certified copies of the priority documents have been received in this National Stage								
application from the International Bureau (PCT Rule 17.2(a)).									
* See the attached detailed Office action for a list of the certified copies not received.									
Attachmen	nt(s)		•	,					
1) 🛛 Notic	ce of References Cited (PTO-892)		4) Interview Summar						
· ==	ce of Draftsperson's Patent Drawing Review (PTO-94 mation Disclosure Statement(s) (PTO/SB/08)	Paper No(s)/Mail E  5) Notice of Informal							
	er No(s)/Mail Date		6)  Other:						

10/814,244 Art Unit: 1797

- 1. Claim 15 is rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention. It is not clear what elements constitute any of the three trademarked spectrometers. Additionally it is not clear if the claim would cover someone that has modified their spectrometer. Additionally, a trademark identifies a manufacturer or a producer, not a product.
- 2. Claims 31-32 provide for the use of a mass spectrometer to analyze a sample containing at least two antiretroviral drugs from at least two classes of antiretroviral drugs, but, since the claim does not set forth any steps involved in the method/process, it is unclear what method/process applicant is intending to encompass. A claim is indefinite where it merely recites a use without any active, positive steps delimiting how this use is actually practiced.
- 3. Claims 31-32 are rejected under 35 U.S.C. 101 because the claimed recitation of a use, without setting forth any steps involved in the process, results in an improper definition of a process, i.e., results in a claim which is not a proper process claim under 35 U.S.C. 101. See for example *Ex parte Dunki*, 153 USPQ 678 (Bd.App. 1967) and *Clinical Products, Ltd.* v. *Brenner*, 255 F. Supp. 131, 149 USPQ 475 (D.D.C. 1966).
- 4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- Claims 1-32 are rejected under 35 U.S.C. 102(b) as being clearly anticipated by Volosov (Clinical Biochemistry March 2002). In the paper Volosov presents a simple rapid method for quantification of antiretrovirals by liquid chromatography-tandem mass spectrometry. In the simple, fast and universal method developed, quantification of any combination of the 15 currently marketed anti-HIV drugs in human plasma is possible, using liquid chromatography-tandem mass spectrometry. The sample preparation section on page 100 teaches that an 80- $\mu$ L plasma sample was spiked with internal standard (cimetidine), and protein was precipitated with 200  $\mu$ L MeCN (acetonitrile). The sample was centrifuged and 30  $\mu$ L aliquot was injected onto the HPLC column, where it underwent an online extraction (sample cleaning) with NH4OAc

Application/Control Number:

10/814,244 Art Unit: 1797

(ammonium acetate, see LC/MS/MS procedure section on page 101). The automatic switching valve was then activated, changing the mobile phase to MeOH and thereby eluting the analytes into the tandem mass spectrometer. Page 100 teaches the mass spectrometer as an API-2000 spectrometer. Stavudine, zidovudine (AZT) and efavirenz were analyzed in the negative ionization mode, while all the other drugs were analyzed in the positive ionization mode (see Table 4). Page 101 teaches that analytes were quantified by multiple reaction monitoring. The high selectivity of a tandem mass analyzer allowed determination of any combination of the drugs within a 4.5-minute run. Between-day precision was <10% for all the analytes at the concentrations tested. Accuracy ranged 95%-105%. The method was linear over the measuring ranges of all the analytes. Within-run precision gave a coefficient of variation of <7% for all the analytes. Good correlation with other analytical methods was observed. The simplicity, universality and high throughput of the method make it suitable for application in a clinical lab. The materials and method section teaches the preparation of standards and serum or plasma quality control specimens.

- 6. Claims 1-2, 4-8, 10, 19, 21-22, 24-26 and 31-32 are rejected under 35 U.S.C. 102(b) as being clearly anticipated by Shoup. In the paper Shoup teaches simultaneous determination of six protease/reverse transcriptase inhibitors in human plasma utilizing LC/MS/MS. With the success of "combination therapies" using reverse transcriptase inhibitors, antiinfectives, and protease inhibitors in the treatment of HIV infection, BAS Analytics developed a single method for profiling six protease/reverse transcriptase inhibitors in human plasma. The method utilizes robotic solid phase extraction at neutral pH and is generally applicable to all the analytes and their internal standards. Page 19 gives the specifics on the tandem mass spectrometer. Page 21 gives data on the standards used.
- 7. The information disclosure statement filed August 15, 2007 fails to comply with 37 CFR 1.98(a)(1), which requires the following: (1) a list of all patents, publications, applications, or other information submitted for consideration by the Office; (2) U.S. patents and U.S. patent application publications listed in a section separately from citations of other documents; (3) the application number of the application in which the information disclosure statement is being submitted on each page of the list; (4) a column that provides a blank space next to each

10/814,244 Art Unit: 1797

document to be considered, for the examiner's initials; and (5) a heading that clearly indicates that the list is an information disclosure statement. The information disclosure statement has been placed in the application file, but the information referred to therein has not been considered unless the reference is listed on a PTO-892 Form.

8. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. The additionally cited art relates to mass spectrometry of compounds, some of which anticipate the instant claims.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Arlen Soderquist whose telephone number is (571) 272-1265. The examiner can normally be reached on Monday-Thursday and Alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jill Warden can be reached on (571) 272-1267. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Arlen Soderquist
Primary Examiner

Art Unit 1797